# Fibrillarin Rabbit mAb

Catalog No.: A27818 Recombinant



#### **Basic Information**

#### **Observed MW**

37KD

#### **Calculated MW**

34kDa

#### Category

SMab Recombinant Monoclonal Antibody

#### **Applications**

WB,IHC-P,IF/ICC,IP,ELISA

### **Cross-Reactivity**

Human, Mouse, Rat

# **Background**

This gene product is a component of a nucleolar small nuclear ribonucleoprotein (snRNP) particle thought to participate in the first step in processing preribosomal RNA. It is associated with the U3, U8, and U13 small nuclear RNAs and is located in the dense fibrillar component (DFC) of the nucleolus. The encoded protein contains an N-terminal repetitive domain that is rich in glycine and arginine residues, like fibrillarins in other species. Its central region resembles an RNA-binding domain and contains an RNP consensus sequence. Antisera from approximately 8% of humans with the autoimmune disease scleroderma recognize fibrillarin.

### **Recommended Dilutions**

WB 1:25000 - 1:100000

**IP** 0.5μg-4μg antibody for 200μg-400μg extracts

of whole cells

**IF/ICC** 1:200 - 1:400

**IHC-P** 1:1000 - 1:4000

**ELISA** Recommended starting

concentration is 1
µg/mL. Please optimize
the concentration
based on your specific

assay requirements.

### **Contact**

www.abclonal.com

## **Immunogen Information**

 Gene ID
 Swiss Prot

 2091
 P22087

#### **Immunogen**

A synthetic peptide corresponding to a sequence with in amino acids 111-210 of human Fibrillarin/U3 RNP (NP\_001427.2).

#### **Synonyms**

FIB; FLRN; Nop1; RNU3IP1

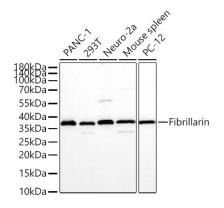
### **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of various lysates using Fibrillarin Rabbit mAb (A27818) at 1:25000 dilution incubated overnight at  $4^{\circ}$ C.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

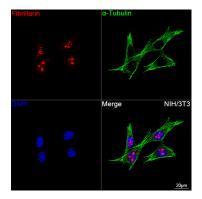
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

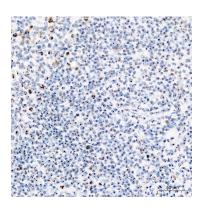
Exposure time: 20s.



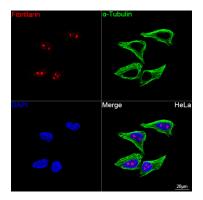
Immunohistochemistry analysis of paraffin-embedded Mouse liver tissue using Fibrillarin Rabbit mAb (A27818) at a dilution of 1:3000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



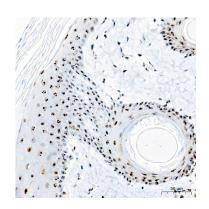
Confocal imaging of NIH/3T3 cells using Fibrillarin Rabbit mAb (A27818, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear



Immunohistochemistry analysis of paraffin-embedded Human tonsil tissue using Fibrillarin Rabbit mAb (A27818) at a dilution of 1:3000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Confocal imaging of HeLa cells using Fibrillarin Rabbit mAb (A27818, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear



Immunohistochemistry analysis of paraffin-embedded Rat skin tissue using Fibrillarin Rabbit mAb (A27818) at a dilution of 1:3000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

10kDa-

staining (Blue). Objective: 100x.

70kDa—

10kDa—

staining (Blue). Objective: 100x.

Immunoprecipitation of Fibrillarin from 300  $\mu$ g extracts of 293T cells was performed using 0.5  $\mu$ g of Fibrillarin Rabbit mAb (A27818). Rabbit IgG isotype control (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1X reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using Fibrillarin Rabbit mAb (A27818) at a dilution of 1:5000.